

## Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area.

Row 1 Administrative Data	Reporter Name <b>Jennifer Greminger</b>	Submission date. <b>October 31, 2018</b>	Contact person (if different than reporter) <b>Joy Thompson</b>	Internal ID <b>32495809</b>
	Address <b>Monsanto Company Mail Stop C3NA 800 N Lindbergh Blvd. St. Louis, MO 63167</b>		Address <b>Missouri Regional Poison Center (MRPC) 7980 Clayton Road, Suite 200 St. Louis, MO 63117</b>	
	Phone # <b>(314) 694-1538</b>		Phone # <b>(314) 772-8300</b>	
	Incident Status: New <input checked="" type="checkbox"/> Update <input type="checkbox"/> If update, include date of original submission.	Location and date of incident. (City, County, State) <b>State: Ohio Date: 9/10/2018</b>	Date registrant became aware of incident. <b>10/17/2018</b>	Was incident part of larger study? Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/>
Row 2  Pesticide(s) Involved	<b>EPA Registration # (Product 1)</b>  <b>71995-33</b>	<b>EPA Registration # (Product 2)</b>	<b>EPA Registration # (Product 3 &amp; 4)</b>	
	A.I. (s) <b>Glyphosate 2% Pelargonic acid 2%</b>	A.I. (s)	A.I. (s)	
	Product 1 Name <b>Roundup Weed and Grass Killer Ready to Use III</b>	Product 2 Name	Product 3&4 Name	
	Exposed to concentrate prior to dilution? Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/> NA <input type="checkbox"/>	Exposed to concentrate prior to dilution? Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/> NA <input type="checkbox"/>	Exposed to concentrate prior to dilution? Y <input type="checkbox"/> N <input type="checkbox"/> U <input type="checkbox"/> NA <input type="checkbox"/>	
	Formulation	Formulation	Formulation	
Row 3  Incident Circumstances	Evidence label directions were not followed? Yes <input type="checkbox"/> No <input type="checkbox"/> U <input checked="" type="checkbox"/> Intentional misuse <input type="checkbox"/>  Applicator certified PCO? Yes <input type="checkbox"/> No <input type="checkbox"/> U <input type="checkbox"/>	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/ woods, agricultural (specify crop) right-of-way (rail, utility, highway).	Situation (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/ formulating). <b>See MRPC incident report (next page)</b>	
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) <b>See MRPC incident report (next page)</b>	Brief description of incident circumstances. <b>See MRPC incident report (next page)</b>  Monsanto asserts that the alleged symptoms are not known effects of this product.		

**Human Exposure / Adverse Effect Incidents  
Involving Monsanto Agricultural Products**

Reporting Categories: H-A, H-B, H-C

Reporting Period: September 1, 2018 – September 30, 2018

<b>Substance:</b>	Roundup Weed and Grass Killer Ready to Use III
<b>Serial Number:</b>	32495809
<b>Date:</b>	9/10/2018
<b>Medical Outcome:</b>	Moderate Effect H-C
<b>EPA Reg. No.</b>	71995-33
<b>Active Ingredients:</b>	Glyphosate 2% Pelargonic acid 2%
<b>State:</b>	Ohio
<b>History and Notes:</b>	69 year old male calling about spraying Roundup Weed and Grass Killer Ready to Use III in early August. He sprayed 2 inches on both side of divider between grass and flowers. He could not eat or walk for 3 weeks. He has lost 22 pounds and is still ill. MRPC discussed the product toxicity. The symptoms do not correlate with the expected response to the product. Advised to continue under care of MD.